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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
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09/716,054 11/17/00 CRABTREE

G STAN-166

HM22/0703

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EXAMINER

COOK, L

ART UNIT

PAPER NUMBER

1641

DATE MAILED:

07/03/01

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trad marks

Office Action Summary	Application No.		Applicant(s)	
	09/716,054		CRABTREE ET AL.	
	Examiner		Art Unit	
	Lisa V. Cook		1641	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In no event, however, may a reply be timely filed SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

1) ☒ Responsive to communication(s) filed on 22 May 2001.

2a) ☐ This action is **FINAL**. 2b) ☐ This action is non-final.

3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

4) ☒ Claim(s) 1-26 is/are pending in the application.

4a) Of the above claim(s) 1-15, 25 and 26 is/are withdrawn from consideration.

5) ☐ Claim(s) _____ is/are allowed.

6) ☒ Claim(s) 16-24 is/are rejected.

7) ☐ Claim(s) _____ is/are objected to.

8) ☒ Claims 1-26 are subject to restriction and/or election requirement.

Application Papers

9) ☐ The specification is objected to by the Examiner.

10) ☐ The drawing(s) filed on _____ is/are objected to by the Examiner.

11) ☐ The proposed drawing correction filed on _____ is: a) ☐ approved b) ☐ disapproved.

12) ☐ The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. § 119

13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) ☐ All b) ☐ Some * c) ☐ None of:

1. ☐ Certified copies of the priority documents have been received.

2. ☐ Certified copies of the priority documents have been received in Application No. _____.

3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

14) ☒ Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).

Attachment(s)

15) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)	18) <input type="checkbox"/> Interview Summary (PTO-413) Paper No(s). _____
16) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)	19) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152)
17) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO-1449) Paper No(s) <u>4</u> .	20) <input type="checkbox"/> Other: _____

DETAILED ACTION

1. Please note that the Examiner of your application in the PTO has changed. To aid in correlating any papers for this application, all correspondence regarding this application should be directed to Group Art Unit **1641**. All communications should be directed to **Lisa V. Cook**, whose telephone number is **(703) 305-0808**.

Election/Restrictions

2. The restriction requirement mailed 4/16/01 (paper#5) erroneously did not include claim 25. Claim 25 being dependent on independent claim 1 should have been included in Group I. Examiner apologizes for the oversight. Applicant's election does not involve claim 25, therefore the correction is presented to clarify the claim status only.

3. Applicant's election with traverse of Group IV (claims 16-24) in Paper No. 10, filed 5/22/01 is acknowledged. Because applicant did not distinctly and specifically point out the supposed errors in the restriction requirement, the election has been treated as an election without traverse (MPEP § 818.03(a)).

The Restriction Requirement is still deemed proper and is therefore made **FINAL**.

4. Currently, claims 1-26 are subject to Restriction and Election Requirement. Claims 1-15 and claims 25-26 have been withdrawn from further consideration by the examiner, 37 CFR 1.142(b), as claims drawn to a non-elected invention. Claims 16-24 are pending and under examination.

Drawings

5. This application has been filed with informal drawings which are acceptable for examination purposes only. Formal drawings will be required when the application is allowed.

Information Disclosure Statement

6. The listing of references in the specification is not a proper information disclosure statement. 37 CFR 1.98(b) requires a list of all patents, publications, or other information submitted for consideration by the Office, and MPEP § 609 A(1) states, "the list may not be incorporated into the specification but must be submitted in a separate paper." Therefore, unless the examiner on form PTO-892 or applicant on form PTO-1449 have cited the references they have not been considered.

7. The information disclosure statement filed 2/12/01 - Paper#4 was considered as to the merits prior to First Action.

Specification

8. The specification has not been checked to the extent necessary to determine the presence of all possible minor errors. Applicant's cooperation is requested in correcting any errors of which applicant may become aware in the specification.

9. The abstract of the disclosure is objected to because page 6, line 35 and page 13, lines 30-33 – list numbers that appear to be U.S. Patent numbers. However the specification does not clearly identify the numbers as such, therefore they could be foreign application numbers, reference identification numbers, etc. Please add "U.S. Patent No." Correction is required. See MPEP § 608.01(b).

Claim Rejections - 35 USC § 112

The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

10. Claims 16-24 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

A. The term "of less than about" in claim 1 is a relative term which renders the claim indefinite. The term "less than about" is not defined by the claim, the specification does not provide a standard for ascertaining the requisite degree, and one of ordinary skill in the art would not be reasonably apprised of the scope of the invention. It is not clear if applicant intends to claim compositions < and = to 5000 Dalton or only compositions <5000 Dalton.

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B. In claim 1, the use of "optionally joined" is indefinite. It is not clear if the limitation is directed to the linking group or the actual linking of the target protein and the blocking protein. In other words are the target protein and the blocking protein always linked to each other (optionally with a linker group or by some other means) or is the limitation to imply that the target protein and blocking protein are not necessarily linked to one another? Please explain.

C. Claim 1 is indefinite because the claim appears to have two different interpretations for the bifunctional inhibitor molecule. In line 4, the bifunctional inhibitor molecule consists of a target protein ligand and a blocking ligand therein being one composition. In line 8, the bifunctional inhibitor molecule, target protein, and blocking protein appear to be separate compositions independent of each other. Is it applicant intent to mean the tripartite complex comprises the bifunctional inhibitor molecule consisting of a target protein and blocking protein further joined to an additional target protein and blocking protein? Please clarify.

D. Claim 1 recites the limitation "an effective amount". The phrase is indefinite when the claim fails to state the function which is to be achieved. Although the claim recites the inhibition of a binding event, because the detection of such inhibition has not been clearly defined by the claims (including essential method steps) the intended function is not known. *In re Frederiksen*, 213 F 2d 547, 102 U.S.P.Q. 35 (C.C.P.A. 1954).

E. Claim 1 is vague and indefinite because it is unclear as to how binding inhibition will occur. Although the claim recites an interaction between a first target protein and a second binding protein in a host, the method does not clearly outline how the second protein and blocking protein interact such that inhibition of the first and second is accomplished. The claims merely read on the formation of a tripartic complex comprising the bifunctional inhibitor molecule, the target protein, and the blocking protein. But does not identify the correlation/interaction/detection allowing for comparative analysis between this tripartic complex and the second binding proteins inhibition. Will the blocking protein and second binding protein compete for the same binding site on the target protein therein allowing for measurement of the blocking protein as an inverse measure for the second binding protein. The method does not including essential method steps.

F. Claim 18 is vague and indefinite because it is not clear whether the bifunctional inhibitor molecule is bound to the target protein and the second binding protein simultaneously or if the bifunctional inhibitor molecule is intended to bind only the second binding protein or the target protein. Further, it is not clear how the bifunctional molecule will simultaneously bind both the second binding protein and target protein at the same site. If the second binding protein is bound in any way to the first target protein how does the bifunctional inhibitor molecule function to inhibit the binding between these two components?

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G. Claim 23 is vague and indefinite in the use of the acronym Hsp90. The term should be defined in its first instance. The initial explanation will convey intended meaning in subsequent abbreviations. Please define.

11. Claims 16-24 are rejected under 35 U.S.C. 112, second paragraph, as being incomplete for omitting essential steps, such omission amounting to a gap between the steps. See MPEP § 2172.01. The omitted steps are elucidated below:

Independent claims 1 is drawn to a method inhibiting a binding event between a first target protein and a second binding protein in a host. Merely reciting a method including the reagents, is not considered to be a proper method (including all the required method steps). Although the specification teaches methods on pages 20-21, the claims do not include the essential method steps. An assay or method, as proposed in the preamble of claims 1, require at least a contact step between reagents and sample, the separation of unbound and bound material, a detection step, and a correlation step. These essential steps for the method have not been outlined for determining protein-protein interaction inhibition. It is suggested that Applicant add steps that at least reflect: (I) a sample and reagent contacting step, (II) the binding or complex formation of a labeled product, the detection of the labeled product, and (III) a correlation step. Please include the necessary steps.

Claim Rejections - 35 USC § 102

12. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

I. Claims 16-24 are rejected under #35 U.S.C. 102(b) as being anticipated by Griffith et al. (Cell, Vol.82, pages 507-522, August 11, 1995).

Griffith et al. disclose the ternary complex of calcineurin A fragment, calcineurin B, FKBP12, and the immunosuppressant drug FK506. The ternary complex provides a structural basis for understanding calcineurin inhibition by FKBP12-FK506. See abstract. The reference meets applicants claimed limitations by teaching the same reagents disclosed in the experimental design present on pages 20-21. The FKBP12-FK506 complex inhibits calcineurin phosphatase activity by blocking the active site from macromolecular phosphorylated substrates like NF-ATp. The information is further taught to be applicable as therapeutic inhibitors. Page 518, Conclusion.

II. Claims 16-21 and 24 are rejected under #35 U.S.C. 102(b) as being anticipated by Varshavsky (Proc. Natl. Acad. Sci. USA Vol.95, pp. 2094-2099, March 1998).

Varshavsky teaches multitarget compounds specific for negative targets concerning the concept of codominant interference. The reference discloses compositions linking two small moiety ligands (< 1Kd page 2095) bipartite compounds consisting of two ligands bound together by a linker (1* and c in Fig. C and D). The ligands are capable of simultaneously binding target protein (C in figure D) and blocking proteins (1 in figure C) thereby possibly forming a tripartite complex. Multitarget drugs designed according to the specific configurations taught by Varshavsky were taught to be useful in the selective killing of cancer cells via the inhibition of a neurotransmitter-inactivating enzyme in a specific subset of the enzyme-containing cells. Therein teaching protein-protein inhibition. See abstract.

Claim Rejections - 35 USC § 103

13. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(f) or (g) prior art under 35 U.S.C. 103(a).

Claims 22 and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Varshavsky (Proc. Natl. Acad. Sci. USA Vol.95, pp. 2094-2099, March 1998) in view of Pouletty et al. (WO 95/10302).

Please see previous discussions of Varshavsky as set forth above.

Varshavsky differs from the instant invention in failing to teach tripartite complexes produced extracellularly.

However, Pouletty et al. teach bifunctional reagents useful in extending in vivo lifetimes of physiologically active agents further reducing the biologically effective concentration or activity of an endogenous or exogenous blood component. Page 2, lines 14-20. A target binding member, which is a physiologically active agent in a mammalian host is bound to a protein via a reagent or conjugate possibly including a linking group. See pages 19 and 20. Applicable proteins include albumin, transferrin, ferritin, and immunoglobulins. See page 3, lines 5-25. The second binding member is usually a macromolecule of at least 5000 Dalton. Page 25, lines 20-25. The bifunctional reagents are taught to have utility in therapeutic methods to detect host derived and foreign targets. Page 5, lines 6-10.

Varshavsky and Poulett et al. are analogous art because they are from the same field of endeavor, both inventions teach techniques involving bifunctional reagents.

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It would have been obvious to one of ordinary skill in the art at the time the invention was made to use the proteins endogenous to the host (i.e. albumin, vitamin receptor, etc..) as taught by Poulett et al. in the method of Varshavsky to perform protein-protein inhibition assay techniques, because such endogenous proteins as taught by Poulett et al. are well known in the art. A person of ordinary skill in the art would have had a reasonable expectation of success utilizing such endogenous proteins, because Poulett et al. taught that the selected blocking protein (long-lived blood component) would affect the manner in which the biological activity of the target is modified and the selection will vary dependent on the nature of the target.

Page 30, lines 23-30. In other words compounds endogenous to the host would cause less side effects and extend dosage levels. Page 1, lines 26-30.

One having ordinary skill in the art would have been motivated to do this because the blocking protein can impart its physiological activities to the target binding member. In this way cellular targets may be inactivated or eliminated. Page 33, lines 16-22.

14. For reasons aforementioned, no claims are allowed.

Remarks

15. Prior art made of record and not relied upon is considered pertinent to the applicant's disclosure:

A. Weiderrecht et al. (U.S. Patent#5,457,182) teach binding interactions involving FK-506 and FKBP12.6.

B. Maragarnore et al. (U.S. Patent#5,242,810) disclose bifunctional inhibitors of platelet activation and thrombin.

16. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform to the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The Group 1641 Fax number is (703) 308-4242, which is able to receive transmissions 24 hours/day, 7 days/week.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Lisa V. Cook whose telephone number is (703) 305-0808. The examiner can normally be reached on Monday-Friday from 8:00 AM - 4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le, can be reached on (703) 305-3399.

Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.


Lisa V. Cook

CM1-7B17

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6/18/01



CHRISTOPHER L. CHIN
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